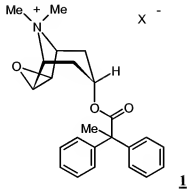
This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

1. (Original) A pharmaceutical composition comprising:

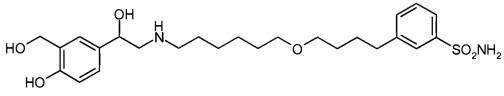
(a) an anticholinergic of formula **1**



1

wherein X^- is an anion with a single negative charge; and

(b) a compound of formula **2**



2,

or the pharmacologically acceptable acid addition salts, solvates, and hydrates thereof.

2. (Original) The pharmaceutical composition according to claim 1, wherein X^- is chloride, bromide, iodide, sulfate, phosphate, methanesulfonate, nitrate, maleate, acetate, citrate, fumarate, tartrate, oxalate, succinate, benzoate, or *p*-toluenesulfonate

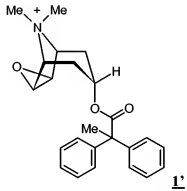
3. (Original) The pharmaceutical composition according to claim 1, further comprising a physiologically acceptable excipient.

4. (Original) The pharmaceutical composition according to claim 1, wherein the compound **2** is a salt of hydrochloric acid, hydrobromic acid, sulfuric acid, phosphoric acid,

methanesulfonic acid, acetic acid, fumaric acid, succinic acid, lactic acid, citric acid, tartaric acid, 1-hydroxy-2-naphthalenecarboxylic acid, or maleic acid.

5. (Currently Amended) The pharmaceutical composition according to claim 1, wherein the compound **2** is an *R*-enantiomer, at least partially separated from the *S*-enantiomer.

6. (Currently Amended) The pharmaceutical composition according to one of claims 1 to 4, wherein the weight ratio of **1'**



1'

to **2** (based on the free base) ~~are~~ is in the range from about 1:30 to about 400:1.

7. (Currently Amended) The pharmaceutical composition according to claim 6, wherein the weight ratio of **1'** to **2** (based on the free base) ~~are~~ is in the range from about 1:25 to about 200:1.

8. (Currently Amended) The pharmaceutical composition according to one of ~~claims 1 to 6~~ claims 1 to 5, wherein the pharmaceutical composition is suitable for inhalation.

9. (Original) The pharmaceutical composition according to claim 8, wherein the pharmaceutical composition is an inhalable powder, propellant-driven metered dose aerosol, or propellant-free inhalable solution.

10. (Original) The pharmaceutical composition according to claim 8, wherein the pharmaceutical composition is an inhalable powder including a physiologically acceptable

excipient selected from monosaccharides, disaccharides, oligo- and polysaccharides, polyalcohols, salts, or mixtures thereof.

11. (Original) The pharmaceutical composition according to one of claims 1, 2, or 5, wherein the pharmaceutical composition is an inhalable powder containing the compounds **1** and **2** as its sole ingredients.

12. (Original) The pharmaceutical composition according to claim 8, wherein the pharmaceutical composition is a propellant-containing inhalable aerosol containing the compounds **1** and **2** in dissolved or dispersed form.

13. (Original) The propellant-containing inhalable aerosol according to claim 12, wherein the propellant gas is a hydrocarbon or halohydrocarbon.

14. (Original) The propellant-containing inhalable aerosol according to claim 12, wherein the propellant gas is *n*-propane, *n*-butane, or isobutane, or a chlorinated and/or fluorinated derivative of methane, ethane, propane, butane, cyclopropane, or cyclobutane, or a mixture thereof.

15. (Original) The propellant-containing inhalable aerosol according to claim 12, wherein the propellant gas is TG11, TG12, TG134a, TG227, or a mixture thereof.

16. (Original) The pharmaceutical composition according to claim 8, wherein the pharmaceutical composition is a propellant-free inhalable solution containing water, ethanol, or a mixture thereof as solvent.

17. (Original) The pharmaceutical composition according to claim 16, further comprising a cosolvent and/or excipient.

18. (New) The pharmaceutical composition according to claim 1, wherein anticholinergics of formula **1** and compounds of formula **2** are substantially the sole active ingredients in the composition.